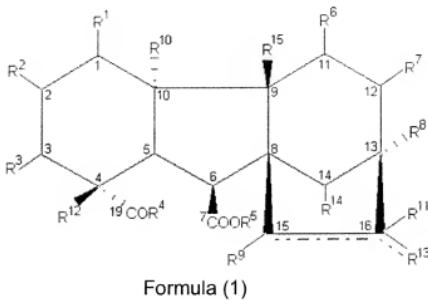


**AMENDMENTS TO THE CLAIMS**

This listing of claims will replace all prior versions and listings of claims in the application:

Claims 1-7: (Canceled)

8. (Currently Amended) A method of treatment for Type II diabetes and its complications and associated conditions comprising administering a compound selected from formula (1) (Gibberellins)



wherein

$R^1$  is H or a group  $-O-R^{20}$ , where  $R^{20}$  is H or a glycosylic ether group (glycoside ether), or  $R^1$  together with  $R^2$  forms a bond (C<sub>1</sub>-C<sub>2</sub> double bond);

$R^2$  is H or a group  $-O-R^{21}$ , where  $R^{21}$  is H, a glycosylic ether group (glycoside ether), or  $R^2$  together with  $R^1$  or  $R^3$  forms a bond (C<sub>1</sub>-C<sub>2</sub> or C<sub>2</sub>-C<sub>3</sub> double bond, respectively);

$R^3$  is H, =O, or  $-O-R^{22}$ , where  $R^{22}$  is H or a glycosylic ether group (glycoside ether), or  $R^3$  together with  $R^2$  forms a bond (C<sub>2</sub>-C<sub>3</sub> double bond);

R<sup>4</sup> together with R<sup>28</sup> forms a bond (lactone);

R<sup>5</sup> is H or a glycosylic ester (glycoside ester) group, or unsubstituted or substituted C<sub>1-20</sub> alkyl esters, allyl esters, active esters;

R<sup>6</sup> is H or OH or together with R<sup>7</sup> forms a bond (C<sub>11</sub>-C<sub>12</sub> double bond);

R<sup>7</sup> is H or -OR<sup>26</sup>, where R<sup>26</sup> is H or a glycosylic ether group (glycoside ether) or R<sup>7</sup> together with R<sup>6</sup> forms a bond (C<sub>11</sub>-C<sub>12</sub> double bond);

R<sup>8</sup> is H[,] or hydroxyl[,] or -OR<sup>27</sup>, where R<sup>27</sup> is a glycosylic ether group (glycoside ether);

R<sup>9</sup> is H or OH;

R<sup>10</sup> is -OR<sup>28</sup>, where R<sup>28</sup> together with R<sup>4</sup> forms a bond (lactone) ;

R<sup>11</sup> is absent;

R<sup>12</sup> is CH<sub>3</sub>;

R<sup>13</sup> is methylene;

R<sup>14</sup> is H;

R<sup>15</sup> is H;

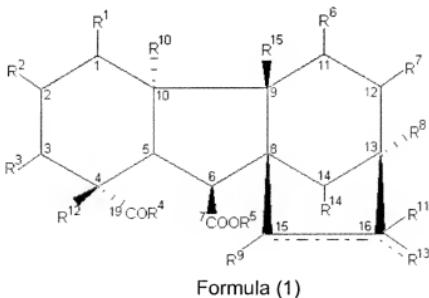
and its pharmaceutically acceptable lactones, esters, active esters, salts and organic bases, in combination with other compatible therapeutic agents selected from the group

consisting of analgesics, anti-hypertensive agents, sedatives, hypnotics, lipid-lowering agents, and anti-infective agents or combinations thereof, to a patient in need thereof.

9. (Previously presented) A method according to claim 11, wherein the Gibberellins are Gibberellin A<sub>3</sub>.

10. (Previously presented) A method according to claim 11, wherein the Gibberellins are a mixture of Gibberellin A<sub>3</sub> and Gibberellin A<sub>4</sub> and/or Gibberellin A<sub>7</sub>.

11. (Currently Amended) A method of treatment for Type I or Type II diabetes and its complications and associated conditions comprising administering compounds selected from formula (1) (Gibberellins)



wherein

R<sup>1</sup> is H or a group -O-R<sup>20</sup>, where R<sup>20</sup> is H or a glycosylic-ether group (glycoside ether), or R<sup>1</sup> together with R<sup>2</sup> forms a bond (C<sub>1</sub>-C<sub>2</sub> double bond);

R<sup>2</sup> is H or a group -O-R<sup>21</sup>, where R<sup>21</sup> is H, a glycosylic-ether group (glycoside ether), or R<sup>2</sup> together with R<sup>1</sup> or R<sup>3</sup> forms a bond (C<sub>1</sub>-C<sub>2</sub> or C<sub>2</sub>-C<sub>3</sub> double bond, respectively);

R<sup>3</sup> is H, =O, or -O-R<sup>22</sup>, where R<sup>22</sup> is H or a glycosylic-ether-group (glycoside-ether), or R<sup>3</sup> together with R<sup>2</sup> forms a bond (C<sub>2</sub>-C<sub>3</sub> double bond);

R<sup>4</sup> together with R<sup>28</sup> forms a bond (lactone);

R<sup>5</sup> is H or a glycosylic-ester (glycoside-ester)-group, or unsubstituted or substituted C<sub>1-20</sub> alkyl esters, allyl esters, active esters;

R<sup>6</sup> is H or OH or together with R<sup>7</sup> forms a bond (C<sub>11</sub>-C<sub>12</sub> double bond);

R<sup>7</sup> is H or -OR<sup>26</sup>, where R<sup>26</sup> is H or a glycosylic-ether-group (glycoside-ether) or R<sup>7</sup> together with R<sup>6</sup> forms a bond (C<sub>11</sub>-C<sub>12</sub> double bond);

R<sup>8</sup> is H[[.]] or hydroxyl[[.]] or -OR<sup>27</sup>, where R<sup>27</sup> is a glycosylic-ether-group (glycoside-ether);

R<sup>9</sup> is H or OH;

R<sup>10</sup> is -OR<sup>28</sup>, where R<sup>28</sup> together with R<sup>4</sup> forms a bond (lactone);

R<sup>11</sup> is absent;

R<sup>12</sup> is CH<sub>3</sub>;

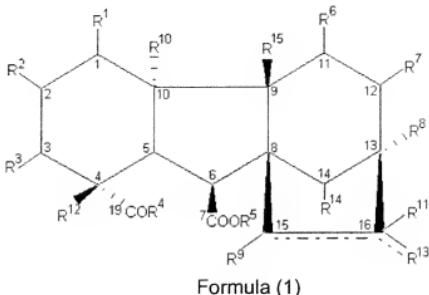
R<sup>13</sup> is methylene;

R<sup>14</sup> is H;

R<sup>15</sup> is H;

and their pharmaceutically acceptable lactones, esters, active esters, salts and organic bases, in combination with substances selected from the group consisting of insulin, its fragment derivatives, IGFs, and growth factors, or combinations thereof, to a patient in need thereof.

12. (Currently Amended) A method of treatment for Type I or Type II diabetes and its complications and associated conditions comprising administering compounds selected from formula (1) (Gibberellins)



wherein

R<sup>1</sup> is H or a group  $-O-R^{20}$ , where R<sup>20</sup> is H or a ~~glycosylic-ether-group (glycoside ether)~~, or R<sup>1</sup> together with R<sup>2</sup> forms a bond (C<sub>1</sub>-C<sub>2</sub> double bond);

R<sup>2</sup> is H or a group  $-O-R^{21}$ , where R<sup>21</sup> is H, a ~~glycosylic-ether-group (glycoside ether)~~, or R<sup>2</sup> together with R<sup>1</sup> or R<sup>3</sup> forms a bond (C<sub>1</sub>-C<sub>2</sub> or C<sub>2</sub>-C<sub>3</sub> double bond, respectively);

R<sup>3</sup> is H, =O, or  $-O-R^{22}$ , where R<sup>22</sup> is H or a ~~glycosylic-ether-group (glycoside ether)~~, or R<sup>3</sup> together with R<sup>2</sup> forms a bond (C<sub>2</sub>-C<sub>3</sub> double bond);

R<sup>4</sup> together with R<sup>28</sup> forms a bond (lactone);

R<sup>5</sup> is H or a glycosylic ester (glycoside ester) group, or unsubstituted or substituted C<sub>1-20</sub> alkyl esters, allyl esters, active esters;

R<sup>6</sup> is H or OH or together with R<sup>7</sup> forms a bond (C<sub>11</sub>-C<sub>12</sub> double bond);

R<sup>7</sup> is H or -OR<sup>26</sup>, where R<sup>26</sup> is H or a glycosylic ether group (glycoside ether) or R<sup>7</sup> together with R<sup>6</sup> forms a bond (C<sub>11</sub>-C<sub>12</sub> double bond);

R<sup>8</sup> is H[[.]] or hydroxyl[[.]] or -OR<sup>27</sup>, where R<sup>27</sup> is a glycosylic ether group (glycoside ether);

R<sup>9</sup> is H or OH;

R<sup>10</sup> is -OR<sup>28</sup>, where R<sup>28</sup> together with R<sup>4</sup> forms a bond (lactone) ;

R<sup>11</sup> is absent;

R<sup>12</sup> is CH<sub>3</sub>;

R<sup>13</sup> is methylene;

R<sup>14</sup> is H;

R<sup>15</sup> is H;

and its pharmaceutically acceptable lactones, esters, active esters, salts and organic bases, in combination with substances selected from the group consisting of insulin, its fragment derivatives, IGFs, and growth factors, or combinations thereof, along with other compatible therapeutic agents selected from the group consisting of analgesics,

anti-hypertensive agents, sedatives, hypnotics, lipid-lowering agents, and anti-infective agents or combinations thereof, to a patient in need thereof.

13. (Previously presented) The method according to claim 11, for the treatment of Type 1 diabetes and its associated conditions.

14. (Previously presented) The method according to claim 11, for the treatment of Type 2 diabetes and its associated conditions.

15. (Previously presented) The method according to claim 14, for the treatment of insulin resistant diabetes.

16. (Currently Amended) The method according to claim 4 11, wherein the diabetic related complications and associated conditions are chosen from obesity, micro and macro vascular diseases, nephropathy, neuropathy and eye diseases.

Claims 17-38: (Canceled)

39. (Previously presented) The method of claim 11, wherein the complications and associated conditions of diabetes are one or more of: obesity, micro and macro vascular diseases, nephropathy, neuropathy, eye diseases, and diabetic ulcerations.

40. (Previously presented) The method of claim 11, wherein the pharmaceutically acceptable salts are selected from alkali metal salts, alkaline earth metal salts, metal, and salts of ammonium or salts of organic bases.

41. (Previously presented) The method of claim 40, wherein the organic bases are lidocaine, or  $NR^{16}R^{17}R^{18}R^{19}$ , where  $R^{16}$ ,  $R^{17}$ ,  $R^{18}$ ,  $R^{19}$ , which may be the same or not the same, are hydrogen, or substituted or unsubstituted  $C_{1-20}$  alkyl, alkanol, or aryl groups.